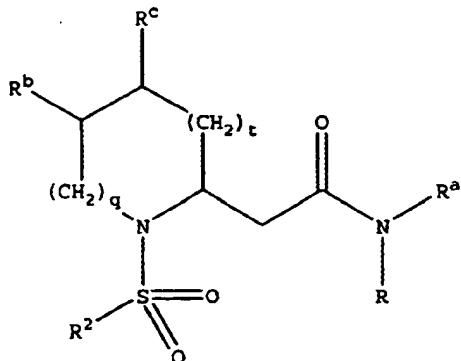


WHAT IS CLAIMED IS:

1. A compound of Formula I'

5



I'

wherein q is 0-3;

wherein t is 0-2, provided that when t is 2, q is not 3;

wherein R is a 9-11 membered fused bicyclic carbocyclic or

10 heterocyclic ring substituted with one to three basic moieties, and optionally substituted with one to three groups independently selected from halo, -NH₂, -OH, -CN, -CF₃, (C₁-C₆)alkylamino, oxo, (C₁-C₆)alkoxy, (C₂-C₆)alkenyl, (C₂-C₆)alkynyl, di(C₁-C₆)alkylamino, -C(O)R⁸, -COOR⁸, -C(O)NR⁸R^{8'}, -NR⁸C(O)R^{8'}, and

15 (C₁-C₆)alkyl, aryl, heteroaryl, cycloalkyl or heterocyclyl, each of which is optionally substituted with one to three groups independently selected from halo, -NH₂, -OH, -CN, -CF₃, (C₁-C₆)alkylamino, halo(C₁-C₆)alkyl, oxo, (C₁-C₆)alkoxy, (C₁-C₆)alkoxy(C₁-C₆)alkyl, (C₁-C₆)alkyl, (C₂-C₆)alkenyl, (C₂-C₆)alkynyl, di(C₁-C₆)alkylamino, -C(O)R⁸, -COOR⁸, -C(O)NR⁸R^{8'}, and -NR⁸C(O)R^{8'};

20 wherein R⁸ and R^{8'} independently are selected from H, and lower alkyl, aryl and heteroaryl, each of which is 25 optionally substituted with one, two or three groups independently selected from lower alkyl,

halogen, lower alkoxy, hydroxy, amino, mono- or dialkylamino, and trifluoromethyl;

wherein R² is selected from arylalkenyl, aryl, and heterocyclyl selected from thienyl, imidazolyl and 5 benzofused heteroaryl, wherein R² is optionally substituted with one to five groups independently selected from halo, -NH₂, -OH, -CN, -CF₃, (C₁-C₆)alkylamino, oxo, (C₁-C₆)alkoxy, haloalkoxy, (C₂-C₆)alkenyl, (C₂-C₆)alkynyl, di(C₁-C₆)alkylamino, -C(O)R⁸, -COOR⁸, -NR⁸C(O)R^{8'}, and 10 (C₁-C₆)alkyl, aryl, heteroaryl, cycloalkyl and heterocyclyl, each of which is optionally substituted with one to three groups independently selected from halo, -NH₂, -OH, -CN, -CF₃, (C₁-C₆)alkylamino, halo(C₁-C₆)alkyl, oxo, (C₁-C₆)alkoxy, (C₁-C₆)alkoxy(C₁-C₆)alkyl, 15 (C₁-C₆)alkyl, (C₂-C₆)alkenyl, (C₂-C₆)alkynyl, di(C₁-C₆)alkylamino, -C(O)R⁸, -COOR⁸, -NR⁸C(O)R^{8'}, and -NR⁸C(O)R^{8''};

wherein R^a is independently selected from H and C₁₋₄-alkyl, 20 and aryl optionally substituted with one to three groups independently selected from halo, -NH₂, -OH, -CN, -CF₃, (C₁-C₆)alkylamino, halo(C₁-C₆)alkyl, oxo, (C₁-C₆)alkoxy, (C₁-C₆)alkoxy(C₁-C₆)alkyl, (C₁-C₆)alkyl, (C₂-C₆)alkenyl, 25 (C₂-C₆)alkynyl, di(C₁-C₆)alkylamino, -C(O)R⁸, -COOR⁸, -C(O)NR⁸R^{8'}, and -NR⁸C(O)R^{8''};

wherein each R^b is independently selected from H, oxo, hydroxy, benzyloxy and C₁₋₂-alkyl;

wherein R^c is independently selected from H and C₁₋₂-alkyl, 30 or wherein R^b and R^c together with the carbon atoms to which they are attached form a 6-membered aryl or heteroaryl ring optionally substituted with one to three groups independently selected from halo, -NH₂, -OH, -CN, -CF₃,

(C₁-C₆)alkylamino, oxo, (C₁-C₆)alkoxy, (C₁-C₆)alkenyl, (C₂-C₆)alkynyl, di(C₁-C₆)alkylamino, -C(O)R^b, -COOR^b, -C(O)NR^bR^b, -NR^bC(O)R^b, and (C₁-C₆)alkyl, aryl, heteroaryl, cycloalkyl and heterocyclyl, each of which is optionally substituted with one to three groups independently selected from halo, -NH₂, -OH, -CN, -CF₃, (C₁-C₆)alkylamino, halo(C₁-C₆)alkyl, oxo, (C₁-C₆)alkoxy, (C₁-C₆)alkoxy(C₁-C₆)alkyl, (C₁-C₆)alkyl, (C₂-C₆)alkenyl, (C₂-C₆)alkynyl, di(C₁-C₆)alkylamino, -C(O)R^b, -COOR^b, -C(O)NR^bR^b, and -NR^bC(O)R^b;

and pharmaceutically acceptable derivatives thereof; provided the basic moiety is not 2-oxo-piperaziny-4-ylmethyl; further provided wherein R^b and R^c do not form a 6-membered aryl when t is 1 and q is 1; further provided the basic substituent is not attached to the bicyclic ring via an oxygen atom; provided R² is not 1-methyimidazol-4-yl; .

20 2. The compound of Claim 1 wherein R is a partially unsaturated carbocyclic ring.

3. The compound of Claim 2 wherein R is 1,2,3,4-tetrahydronaphthyl.

25 4. The compound of Claim 2 wherein R is indanyl.

5. The compound of Claim 2 wherein R is selected from 1,2,3,4-tetrahydronaphth-1-yl, 1,2,3,4-tetrahydronaphth-2-yl, indan-1-yl and indan-2-yl.

30 6. The compound of Claim 1 wherein R is partially unsaturated heterocyclyl.

7. The compound of Claim 6 wherein R is chroman.

8. The compound of Claim 6 wherein R is 2,2-dioxo-3,4-dihydro-1H-2,1-benzothiazinyl.

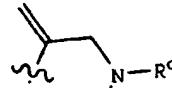
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9. The compound of Claim 1 wherein R is chroman-4-yl, 5,6,7,8-tetrahydro-quinazolin-5-yl, 5,6,7,8-tetrahydro-[1,6]naphthyridin-4-yl or 2,2-dioxo-3,4-dihydro-1H-2,1-benzothiazin-4-yl.

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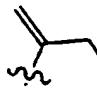
10. The compound of Claim 1 wherein
q is 1 or 2;
t is 0 or 1;
wherein each R² is selected from phenyl-CH=CH-,
15 tetrahydronaphthyl, naphtho[2.3-d]dioxolyl, benzofuranyl, benzoxadiazolyl, benzothiadiazolyl, benzothiazolyl, 1H-pyrazolyl, thienyl, isoxazolthienyl, benzothienyl, thieno[3,2-c]pyridinyl, naphthyl, phenyl, pyridinyl, tetrahydroisoquinolyl, quinolinyl and isoquinolinyl;
20 wherein R² is optionally substituted with one to five groups independently selected from halo, -NH₂, -OH, -CN, -CF₃, (C₁-C₆)alkylamino, oxo, (C₁-C₆)alkoxy, (C₂-C₆)alkenyl, (C₂-C₆)alkynyl, di(C₁-C₆)alkylamino, -C(O)R⁸, -COOR⁸, -C(O)NR⁸R⁸', -NR⁸C(O)R⁸', and
25 (C₁-C₆)alkyl, aryl, heteroaryl, cycloalkyl or heterocyclyl, each of which is optionally substituted with one to three groups independently selected from halo, -NH₂, -OH, -CN, -CF₃, (C₁-C₆)alkylamino, halo(C₁-C₆)alkyl, oxo, (C₁-C₆)alkoxy, (C₁-C₆)alkoxy(C₁-C₆)alkyl,
30 (C₁-C₆)alkyl, (C₂-C₆)alkenyl, (C₂-C₆)alkynyl, di(C₁-C₆)alkylamino, -C(O)R⁸, -COOR⁸, -C(O)NR⁸R⁸', and -NR⁸C(O)R⁸';
wherein R^a is selected from H and C₁₋₂-alkyl;
wherein R^b and R^c are H;

wherein the basic substituent on R is selected from
 cycloalkylamino(C₁-C₆)alkyl, cycloalkyl(C₁-



C₆)alkylamino(C₁-C₆)alkyl, R^e, heteroarylamino(C₁-
 C₆)alkyl, heteroaryl(C₁-C₆)alkylamino(C₁-C₆)alkyl,
 5 arylamino(C₁-C₆)alkyl, alkoxyalkylaminoalkyl,
 hydroxyalkylaminoalkyl, alkenylalkylaminoalkyl,
 aminocarbonylalkylamino-alkyl, carboxyalkylaminoalkyl,
 aryl(C₁-C₆)alkylamino(C₁-C₆)alkyl, C₁-C₆-alkylamino-C₁-C₆-
 alkoxy, haloalkylaminoalkyl, amino(C₁-C₆)alkyl, (C₁-
 10 C₆)alkylamino(C₁-C₆)alkyl, 5-8 membered nitrogen-
 containing heterocyclyl, 5-7 membered nitrogen-containing
 heterocyclyl-alkylaminoalkyl and 5-7 membered
 heterocyclyl-alkyl; and wherein each of said basic
 substituents is optionally substituted with one to three
 15 groups independently selected from halo, -NH₂, -OH, -CN,
 -CF₃, (C₁-C₆)alkylamino, oxo, (C₁-C₆)alkoxy, (C₂-
 C₆)alkenyl, (C₂-C₆)alkynyl, di(C₁-C₆)alkylamino, -C(O)R⁸, -
 COOR⁸, -C(O)NR⁸R⁸', -NR⁸C(O)R⁸', and
 (C₁-C₆)alkyl, aryl, heteroaryl, cycloalkyl or
 20 heterocyclyl, each of which is optionally substituted
 with one to three groups independently selected from
 halo, -NH₂, -OH, -CN, -CF₃, (C₁-C₆)alkylamino, halo(C₁-
 C₆)alkyl, oxo, (C₁-C₆)alkoxy, (C₁-C₆)alkoxy(C₁-C₆)alkyl,
 (C₁-C₆)alkyl, (C₂-C₆)alkenyl, (C₂-C₆)alkynyl, di(C₁-
 25 C₆)alkylamino, -C(O)R⁸, -COOR⁸, -C(O)NR⁸R⁸', and -
 NR⁸C(O)R⁸'; and
 wherein R^d is selected from alkyl, cycloalkyl,
 cycloalkylalkyl, hydroxyalkyl, alkoxyalkyl, and H;
 wherein R^e is H; or where R^d and R^e together with the
 30 nitrogen atom to which they are attached form a
 heterocyclic ring;
 and pharmaceutically acceptable derivatives thereof.

11. The compound of Claim 1 wherein R² is selected from phenyl-CH=CH-, tetrahydronaphthyl, naphtho[2.3-d]dioxol-6-yl, 1-benzofur-2-yl, 2,1,3-benzoxadiazol-4-yl, 2,1,3-5 benzothiadiazol-4-yl, 1,3-benzothiazol-2-yl, 1H-pyrazol-4-yl, thien-2-yl, 5-isoxazolthien-2-yl, benzothien-2-yl, benzothien-3-yl, thieno[3,2-c]pyridin-2-yl, naphthyl, phenyl, 3-pyridyl, tetrahydroisoquinolyl, quinol-8-yl and isoquinolyl; wherein each R² is said optionally substituted; 10 wherein R^a is H; and wherein the basic substituent on R is selected from -NH₂,



, C₃₋₆-cycloalkyl(C_{1-C₂})alkylamino(C_{1-C₂})alkyl, C₃₋₆-cycloalkylamino(C_{1-C₂})alkyl, (C_{1-C₂})alkoxy(C_{1-C₂})alkylamino(C_{1-C₂})alkyl, mono-C₂₋₄-alkenylamino-C₁₋₄-alkyl, di-C₂₋₄-alkenylamino-C₁₋₄-alkyl, hydroxy-C₁₋₄-alkylamino-C₁₋₄-alkyl, aminocarbonyl-C₁₋₄-alkylamino-C₁₋₂-alkyl, mono-C₁₋₆-alkylamino-C₁₋₄-alkyl, di-C₁₋₄-alkylamino-C₁₋₄-alkyl and 5-8 membered heterocyclyl-C₁₋₄-alkyl; 15 wherein each is optionally substituted with one to three groups independently selected from halo, -NH₂, -OH, -CN, -CF₃, (C_{1-C₆})alkylamino, oxo, (C_{1-C₆})alkoxy, (C_{2-C₆})alkenyl, (C_{2-C₆})alkynyl, di(C_{1-C₆})alkylamino, -C(O)R⁸, -COOR⁸, -C(O)NR⁹R^{8'}, -NR⁹C(O)R^{8'}, and (C_{1-C₆})alkyl, aryl, heteroaryl, 20 cycloalkyl or heterocyclyl, each of which is optionally substituted with one to three groups independently selected from halo, -NH₂, -OH, -CN, -CF₃, (C_{1-C₆})alkylamino, halo(C_{1-C₆})alkyl, oxo, (C_{1-C₆})alkoxy, (C_{1-C₆})alkoxy(C_{1-C₆})alkyl, (C_{1-C₆})alkyl, (C_{2-C₆})alkenyl, (C_{2-C₆})alkynyl, di(C_{1-C₆})alkylamino, -C(O)R⁸, -COOR⁸, -C(O)NR⁹R^{8'}, and -NR⁹C(O)R^{8'}; 25 30

wherein R^d is selected from C₁₋₅-alkyl, C₃₋₆-cycloalkyl, C₃₋₆-cycloalkyl-C₁₋₄-alkyl, C₁₋₄-hydroxyalkyl, C₁₋₃-alkoxy-C₁₋₃-alkyl and H; and

wherein R^e is H; or where R^d and R^e together with the
5 nitrogen atom to which they are attached form a 4-8
membered nitrogen-containing heterocyclic ring;
and pharmaceutically acceptable derivatives thereof.

12. The compound of Claim 1 wherein R^a is H.

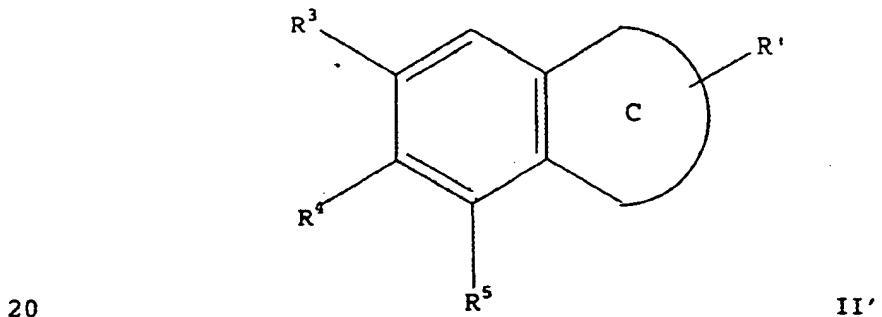
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13. The compound of Claim 1 wherein the basic
substituent on R is selected from -NH₂, aminomethyl,
aminoethyl, aminopropyl, isopropylaminomethyl, t-
butylaminomethyl, iso-butylaminomethyl, 1-
15 methylpropylaminomethyl, 2-methylbutylaminomethyl, 2,2'-
dimethylpropylaminomethyl, 2,2',3-trimethylpropylaminomethyl,
allyl-aminomethyl, isopropylaminopropyl, 1-
(isobutylamino)ethyl, 1-(isopropylamino)-1-methylethyl, N-
isopropyl-N-ethylaminomethyl, N-isopropyl-N-
20 methylaminomethyl, N-t-butyl-N-methylaminomethyl, N-iso-
butyl-N-methylaminomethyl, N-t-butyl-N-ethylaminomethyl, N-
isobutyl-N-methylaminomethyl, N-t-butyl-N-
isopropylaminomethyl, N,N-di(isopropyl)aminomethyl, N,N-
dimethylaminomethyl, N,N-diethylaminomethyl, N,N-di(t-
25 butyl)-aminomethyl, N,N-di(allyl)-aminomethyl,
cyclopropylaminomethyl, 1-(cyclopropylamino)ethyl,
cyclobutylaminomethyl, 2-(cyclobutylamino)ethyl, 1-
(cyclobutylamino)ethyl, cyclopentylaminomethyl, 1-
cyclopentylaminoethyl, cyclopropylmethylaminomethyl,
30 hydroxyethylamino-allyl, isopropylamino-allyl, t-butylamino-
allyl, cyclopropylmethylamino-allyl, piperidin-1-yl-allyl,
pyrrolidin-1-yl-allyl, azetidin-1-yl-allyl, 3-
hydroxypyrrolidin-1-yl-allyl, aminocarbonylethylaminomethyl,
methoxyethylaminomethyl, 1-(methoxyethylamino)ethyl, 1-

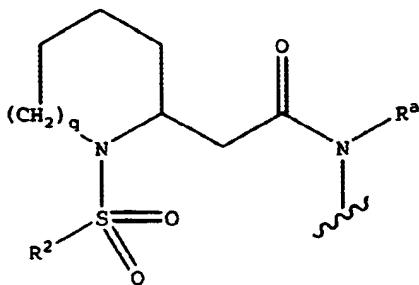
piperidinylmethyl, 2-(piperidin-1-yl)ethyl, 3,4-dihydropiperidin-1-ylmethyl, 4-fluoropiperidinylmethyl, 4,4'-difluoropiperidinylmethyl, 4-(piperidin-1-yl)piperidinylmethyl, 3-aminocarbonylpiperidin-1-ylmethyl,
 5 4-(dimethylamino)piperidin-1-ylmethyl, 2,6-dimethylpiperidin-1-ylmethyl, 3,3-dimethylpiperidin-1-ylmethyl, piperidin-1-yl-2-methylethyl, 3-hydroxypiperidin-1-yl, 4-morpholinylmethyl, 4-morpholinylethyl, 1-pyrrolidinylmethyl, 2-methylpyrrolidin-1-ylmethyl, 1-(methylpyrrolidin-1-yl)ethyl, 2,5-dimethylpyrrolidin-1-ylmethyl, 1-azetidinylmethyl, 7-aza-bicyclo[2.2.1]heptyl, 10 piperazin-1-ylmethyl, 4-methylpiperazin-1-ylmethyl, and 1-pyrrolidinylethylaminomethyl.

15 14. The compound of Claim 1 wherein R^b and R^c are joined to form a phenyl ring; and wherein q is 2.

15. A compound of Formula II'



wherein the C ring is a 4- to 7- membered saturated carbocyclic or heterocyclic moiety;
 wherein R' is selected from



wherein q is 0-3;

wherein R^2 is selected from arylalkenyl, aryl, and

5 heterocyclyl selected from thienyl, imidazolyl and
benzofused heteroaryl, wherein R^2 is optionally
substituted with one to five groups independently
selected from halo, $-NH_2$, $-OH$, $-CN$, $-CF_3$, (C_1-C_6) alkylamino, oxo, (C_1-C_6) alkoxy, haloalkoxy, (C_2-C_6) alkenyl, (C_2-C_6) alkynyl, di (C_1-C_6) alkylamino, $-C(O)R^8$, $-COOR^8$, $-C(O)NR^8R^8'$, $-NR^8C(O)R^8'$, and

10 (C_1-C_6) alkyl, aryl, heteroaryl, cycloalkyl and
heterocyclyl, each of which is optionally substituted
with one to three groups independently selected from
halo, $-NH_2$, $-OH$, $-CN$, $-CF_3$, (C_1-C_6) alkylamino, halo (C_1-C_6) alkyl, oxo, (C_1-C_6) alkoxy, (C_1-C_6) alkoxy (C_1-C_6) alkyl,
15 (C_1-C_6) alkyl, (C_2-C_6) alkenyl, (C_2-C_6) alkynyl, di (C_1-C_6) alkylamino, $-C(O)R^8$, $-COOR^8$, $-C(O)NR^8R^8'$, and $-NR^8C(O)R^8'$;

20 wherein R^a is independently selected from H and C_{1-4} -alkyl,
or
aryl optionally substituted with one to three groups
selected from halo, $-NH_2$, $-OH$, $-CN$, $-CF_3$, (C_1-C_6) alkylamino, halo (C_1-C_6) alkyl, oxo, (C_1-C_6) alkoxy,
25 (C_1-C_6) alkoxy (C_1-C_6) alkyl, (C_1-C_6) alkyl, (C_2-C_6) alkenyl,
 (C_2-C_6) alkynyl, di (C_1-C_6) alkylamino, $-C(O)R^8$, $-COOR^8$,
 $-C(O)NR^8R^8'$, and $-NR^8C(O)R^8'$;

wherein R³, R⁴ and R⁵ are the same or different and represent
H, halo, -NH₂, -OH, -CN, -CF₃, (C₁-C₆)alkylamino, oxo, (C₁-
C₆)alkoxy, (C₂-C₆)alkenyl, (C₂-C₆)alkynyl, di(C₁-
C₆)alkylamino, -C(O)R⁸, -COOR⁸, -C(O)NR⁸R^{8'}, -NR⁸C(O)R^{8''}, a
5 basic moiety, or
(C₁-C₂)alkyl, aryl, heteroaryl, cycloalkyl or
heterocyclyl, each of which is optionally substituted
with one to three groups independently selected from
halo, -NH₂, -OH, -CN, -CF₃, (C₁-C₆)alkylamino, halo(C₁-
C₆)alkyl, oxo, (C₁-C₆)alkoxy, (C₁-C₆)alkoxy(C₁-C₆)alkyl,
10 (C₁-C₆)alkyl, (C₂-C₆)alkenyl, (C₂-C₆)alkynyl, di(C₁-
C₆)alkylamino, -C(O)R⁸, -COOR⁸, -C(O)NR⁸R^{8'}, and -
NR⁸C(O)R^{8''}; and
wherein R⁸ and R^{8'} independently are selected from H, and
15 lower alkyl, aryl and heteroaryl, each of which is
optionally substituted with one, two or three groups
independently selected from lower alkyl, halogen,
lower alkoxy, hydroxy, amino, mono- or dialkylamino,
and trifluoromethyl;
20 provided at least one of R³, R⁴ and R⁵ is a basic moiety;
and pharmaceutically acceptable derivatives thereof.

16. The compound of Claim 15 wherein R³ and R⁵ are H; and
wherein R⁴ is selected from -NH₂, aminomethyl, aminoethyl,
25 aminopropyl, isopropylaminomethyl, t-butylaminomethyl, iso-
butylaminomethyl, 1-methylpropylaminomethyl, 2-
methylbutylaminomethyl, 2,2'-dimethylpropylaminomethyl,
2,2',3-trimethylpropylaminomethyl, allyl-aminomethyl,
isopropylaminopropyl, 1-(isobutylamino)ethyl, 1-
30 (isopropylamino)-1-methylethyl, N-isopropyl-N-
ethylaminomethyl, N-isopropyl-N-methylaminomethyl, N-t-
butyl-N-methylaminomethyl, N-iso-butyl-N-methylaminomethyl,
N-t-butyl-N-ethylaminomethyl, N-isobutyl-N-
methylaminomethyl, N-t-butyl-N-isopropylaminomethyl, N,N-

di(isopropyl)aminomethyl, N,N-dimethylaminomethyl, N,N-diethylaminomethyl, N,N-di(*t*-butyl)-aminomethyl, N,N-di(allyl)-aminomethyl, cyclopropylaminomethyl, 1-(cyclopropylamino)ethyl, cyclobutylaminomethyl, 2-(cyclobutylamino)ethyl, 1-(cyclobutylamino)ethyl, cyclopentylaminomethyl, 1-cyclopentylaminoethyl, cyclopropylmethylaminomethyl, hydroxyethylamino-allyl, isopropylamino-allyl, *t*-butylamino-allyl, cyclopropylmethylamino-allyl, piperidin-1-yl-allyl,
10 pyrrolidin-1-yl-allyl, azetidin-1-yl-allyl, 3-hydroxypyrrolidin-1-yl-allyl, aminocarbonylethylaminomethyl, methoxyethylaminomethyl, 1-(methoxyethylamino)ethyl, 1-piperidinylmethyl, 2-(piperidin-1-yl)ethyl, 3,4-dihydropiperidin-1-ylmethyl, 4-fluoropiperidinylmethyl,
15 4,4'-difluoropiperidinylmethyl, 4-(piperidin-1-yl)piperidinylmethyl, 3-aminocarbonylpiperidin-1-ylmethyl, 4-(dimethylamino)piperidin-1-ylmethyl, 2,6-dimethylpiperidin-1-ylmethyl, 3,3-dimethylpiperidin-1-ylmethyl, piperidin-1-yl-2-methylethyl, 3-hydroxypiperidin-1-yl, 4-morpholinylmethyl, 4-morpholinyethyl, 1-pyrrolidinylmethyl, 2-methylpyrrolidin-1-ylmethyl, 1-(methylpyrrolidin-1-yl)ethyl, 2,5-dimethylpyrrolidin-1-ylmethyl, 1-azetidinylmethyl, 7-aza-bicyclo[2.2.1]heptyl, piperazin-1-ylmethyl, 4-methylpiperazin-1-ylmethyl, and 1-pyrrolidinylethylaminomethyl;
25 and pharmaceutically acceptable derivatives thereof.

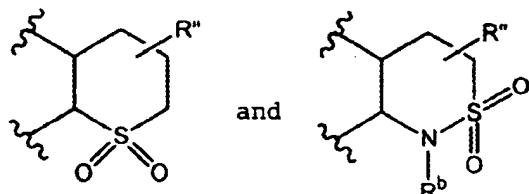
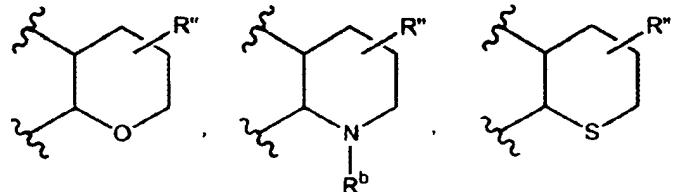
17. The compound of Claim 15 wherein R¹ and R⁵ are H; and wherein R³ is selected from -NH₂, aminomethyl, 30 aminoethyl, aminopropyl, isopropylaminomethyl, *t*-butylaminomethyl, iso-butylaminomethyl, 1-methylpropylaminomethyl, 2-methylbutylaminomethyl, 2,2'-dimethylpropylaminomethyl, 2,2',3-trimethylpropylaminomethyl, allyl-aminomethyl,

isopropylaminopropyl, 1-(isobutylamino)ethyl, 1-(isopropylamino)-1-methylethyl, N-isopropyl-N-ethylaminomethyl, N-isopropyl-N-methylaminomethyl, N-t-butyl-N-methylaminomethyl, N-iso-butyl-N-methylaminomethyl,
5 N-t-butyl-N-ethylaminomethyl, N-isobutyl-N-methylaminomethyl, N-t-butyl-N-isopropylaminomethyl, N,N-di(isopropyl)aminomethyl, N,N-dimethylaminomethyl, N,N-diethylaminomethyl, N,N-di(t-butyl)-aminomethyl, N,N-di(allyl)-aminomethyl, cyclopropylaminomethyl, 1-(cyclopropylamino)ethyl, cyclobutylaminomethyl, 2-(cyclobutylamino)ethyl, 1-(cyclobutylamino)ethyl, cyclopentylaminomethyl, 1-cyclopentylaminoethyl, cyclopropylmethylaminomethyl, hydroxyethylamino-allyl, isopropylamino-allyl, t-butylamino-allyl,
10 15 cyclopropylmethylamino-allyl, piperidin-1-yl-allyl, pyrrolidin-1-yl-allyl, azetidin-1-yl-allyl, 3-hydroxypyrrolidin-1-yl-allyl, aminocarbonylethylaminomethyl, methoxyethylaminomethyl, 1-(methoxyethylamino)ethyl, 1-piperidinylmethyl, 2-(piperidin-1-yl)ethyl, 3,4-dihydropiperidin-1-ylmethyl, 4-fluoropiperidinylmethyl, 4,4'-difluoropiperidinylmethyl, 4-(piperidin-1-yl)piperidinylmethyl, 3-aminocarbonylpiperidin-1-ylmethyl, 4-(dimethylamino)piperidin-1-ylmethyl, 2,6-dimethylpiperidin-1-ylmethyl, 3,3-dimethylpiperidin-1-ylmethyl, piperidin-1-yl-2-methylethyl, 3-hydroxypiperidin-1-yl, 4-morpholinylmethyl, 4-morpholinylethyl, 1-pyrrolidinylmethyl, 2-methylpyrrolidin-1-ylmethyl, 1-(methylpyrrolidin-1-yl)ethyl, 2,5-dimethylpyrrolidin-1-ylmethyl, 1-azetidinylmethyl, 7-aza-bicyclo[2.2.1]heptyl, piperazin-1-ylmethyl, 4-methylpiperazin-1-ylmethyl, and 1-pyrrolidinylethylaminomethyl;
30 and pharmaceutically acceptable derivatives thereof.

18. The compound of Claim 15 wherein R³ and R⁴ are H; and wherein R⁵ is selected from -NH₂, aminomethyl, aminoethyl, aminopropyl, isopropylaminomethyl, t-butylaminomethyl, iso-butylaminomethyl, 1-
5 methylpropylaminomethyl, 2-methylbutylaminomethyl, 2,2'-dimethylpropylaminomethyl, 2,2',3-trimethylpropylaminomethyl, allyl-aminomethyl, isopropylaminopropyl, 1-(isobutylamino)ethyl, 1-(isopropylamino)-1-methylethyl, N-isopropyl-N-
10 methylaminomethyl, N-t-butyl-N-methylaminomethyl, N-iso-butyl-N-methylaminomethyl, N-t-butyl-N-ethylaminomethyl, N-isobutyl-N-methylaminomethyl, N-t-butyl-N-isopropylaminomethyl, N,N-di(isopropyl)aminomethyl, N,N-dimethylaminomethyl, N,N-diethylaminomethyl, N,N-di(t-
15 butyl)-aminomethyl, N,N-di(allyl)-aminomethyl, cyclopropylaminomethyl, 1-(cyclopropylamino)ethyl, cyclobutylaminomethyl, 2-(cyclobutylamino)ethyl, 1-(cyclobutylamino)ethyl, cyclopentylaminomethyl, 1-cyclopentylaminoethyl, cyclopropylmethylaminomethyl,
20 hydroxyethylamino-allyl, isopropylamino-allyl, t-butylamino-allyl, cyclopropylmethylamino-allyl, piperidin-1-yl-allyl, pyrrolidin-1-yl-allyl, azetidin-1-yl-allyl, 3-hydroxypyrrolidin-1-yl-allyl, aminocarbonylethylaminomethyl, methoxyethylaminomethyl, 1-(methoxyethylamino)ethyl, 1-
25 piperidinylmethyl, 2-(piperidin-1-yl)ethyl, 3,4-dihdropiperidin-1-ylmethyl, 4-fluoropiperidinylmethyl, 4,4'-difluoropiperidinylmethyl, 4-(piperidin-1-yl)piperidinylmethyl, 3-aminocarbonylpiperidin-1-ylmethyl, 4-(dimethylamino)piperidin-1-ylmethyl, 2,6-
30 dimethylpiperidin-1-ylmethyl, 3,3-dimethylpiperidin-1-ylmethyl, piperidin-1-yl-2-methylethyl, 3-hydroxypiperidin-1-yl, 4-morpholinylmethyl, 4-morpholinylethyl, 1-pyrrolidinylmethyl, 2-methylpyrrolidin-1-ylmethyl, 1-(methylpyrrolidin-1-yl)ethyl, 2,5-dimethylpyrrolidin-1-

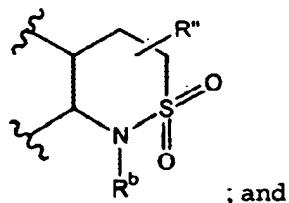
ylmethyl, 1-azetidinylmethyl, 7-aza-bicyclo[2.2.1]heptyl, piperazin-1-ylmethyl, 4-methylpiperazin-1-ylmethyl, and 1-pyrrolidinylethylaminomethyl.

5 19. The compound of Claim 15 wherein the C ring is selected from



10 wherein R^b is independently selected from R' , H and C_{1-2} -alkyl; and wherein R'' is R' when R^b is hydrogen or C_{1-2} -alkyl, or R'' is hydrogen when R^b is R' .

15 20. The compound of Claim 19 wherein the C ring is



wherein R^b is R' .

21. The compound of Claim 15 wherein R² is selected from phenyl-CH=CH-, tetrahydronaphthyl, naphtho[2.3-d]dioxol-6-yl, 1-benzofuran-2-yl, 2,1,3-benzoxadiazol-4-yl, 2,1,3-benzothiadiazol-4-yl, 1,3-benzothiazol-2-yl, 1H-pyrazol-4-yl, thien-2-yl, 5-isoxazolthien-2-yl, benzothien-2-yl, benzothien-3-yl, thieno[3,2-c]pyridin-2-yl, naphthyl, phenyl, 3-pyridyl, tetrahydroisoquinolinyl, quinolinyl and isoquinolinyl;

wherein each R² is optionally substituted with one to five groups independently selected from halo, -NH₂, -OH, -CN, -CF₃, (C₁-C₆)alkylamino, oxo, (C₁-C₆)alkoxy, (C₂-C₆)alkenyl, (C₂-C₆)alkynyl, di(C₁-C₆)alkylamino, -C(O)R⁸, -COOR⁸, -C(O)NR⁸R⁸', -NR⁸C(O)R⁸', and (C₁-C₆)alkyl, aryl, heteroaryl, cycloalkyl and heterocyclyl, each of which is optionally substituted with one to three groups independently selected from halo, -NH₂, -OH, -CN, -CF₃, (C₁-C₆)alkylamino, halo(C₁-C₆)alkyl, oxo, (C₁-C₆)alkoxy, (C₁-C₆)alkoxy(C₁-C₆)alkyl, (C₁-C₆)alkyl, (C₂-C₆)alkenyl, (C₂-C₆)alkynyl, di(C₁-C₆)alkylamino, -C(O)R⁸, -COOR⁸, -C(O)NR⁸R⁸', or -NR⁸C(O)R⁸'.

22. The compound of Claim 15 wherein R² is selected from 2-naphthyl, 1-naphthyl, phenyl, 3-chlorophenyl, 4-chlorophenyl, 3,5-dichlorophenyl, 3,4-dichlorophenyl, 2,4,6-trichlorophenyl, 3-fluorophenyl, 3-methoxyphenyl, 4-methoxyphenyl, 3-biphenyl, 4'-chlorophenyl-3-phenyl, 3-methylphenyl, 3-trifluoromethylphenyl, 2-chlorobenzothien-3-yl, and 3-pyridyl; wherein R² is optionally substituted with one or more groups selected from halo, -NH₂, -OH, -CO₂H, (C₁-C₂)alkylamino, (C₁-C₂)alkoxy, (C₁-C₂)alkoxy-(C₁-C₂)alkyl, (C₁-C₂)alkyl, halo(C₁-C₂)alkyl, di(C₁-C₂)alkylamino, and phenyl.

23. The compound of Claim 15 wherein R⁸ is H.

24. The compound of Claim 15 wherein R^2 is 2-naphthyl.

25. The compound of Claim 15 wherein R² is 3,4-
5 dichlorophenyl.

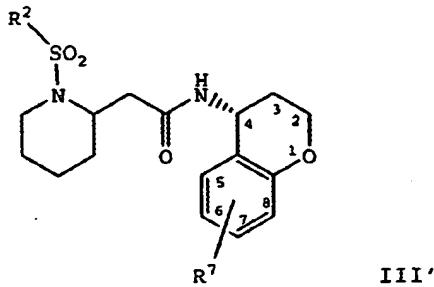
26. The compound of Claim 15 wherein R² is 3-trifluoromethylphenyl.

10 27. The compound of Claim 1 and/or pharmaceutically acceptable derivatives thereof selected from

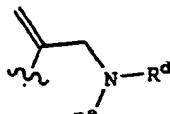
N-(7-Piperidin-1-ylmethyl-chroman-4-(R)-yl)-2-[1-(3-trifluoromethyl-benzenesulfonyl)-piperidin-2-yl]-acetamide; 2-[1-(Naphthalene-2-sulfonyl)-piperidin-2-yl]-N-(7-piperidin-1-ylmethyl-chroman-4-(R)-yl)-acetamide; and 2-[1-(Naphthalene-2-sulfonyl)-pyrrolidin-2-(L)-yl]-N-(7-piperidin-1-ylmethyl-chroman-4-(R)-yl)-acetamide.

20

28. A compound of Formula III'



25 wherein R² is selected from naphthyl, phenyl, pyridinyl, benzothienyl, quinolinyl and isoquinolinyl, and wherein each is optionally substituted with one to three substituents selected from chloro, fluoro, methoxy, methyl, trifluoromethyl, and phenyl; and



wherein R' is selected from R^e , C_{3-6} -cycloalkyl (C_1-C_2)alkylamino (C_1-C_2)alkyl, C_{3-6} -cycloalkylamino (C_1-C_2)alkyl, (C_1-C_2)alkoxy (C_1-C_2)alkylamino (C_1-C_2)alkyl, mono- C_{2-4} -alkenylamino- C_{1-4} -alkyl, di- C_{2-4} -alkenylamino- C_{1-4} -alkyl, 5-8 membered heterocyclyl- C_{1-4} -alkyl; wherein the 5-8 membered heterocyclyl- $(\text{CH}_2)_p$ is optionally substituted with one to three groups independently selected from halo, $-\text{NH}_2$, $-\text{OH}$, $-\text{CN}$, $-\text{CF}_3$, (C_1-C_6)alkylamino, oxo, (C_1-C_6)alkoxy, (C_2-C_6)alkenyl, (C_2-C_6)alkynyl, di(C_1-C_6)alkylamino, $-\text{C}(\text{O})\text{R}^8$, $-\text{COOR}^8$, $-\text{C}(\text{O})\text{NR}^8\text{R}^{8'}$, $-\text{NR}^8\text{C}(\text{O})\text{R}^{8'}$, $=\text{NCN}$;

wherein R^d is selected from C_{1-5} -alkyl, C_{3-6} -cycloalkyl, C_{3-6} -cycloalkyl- C_{1-4} -alkyl, C_{1-4} -hydroxyalkyl, C_{1-3} -alkoxy- C_{1-3} -alkyl and H; and

wherein R^e is H; or where R^d and R^e together with the nitrogen atom to which they are attached form a 4-8 membered nitrogen-containing heterocyclic ring;

wherein R' is at position 6, 7 or 8; and

wherein R^8 and $\text{R}^{8'}$ independently are selected from H, and lower alkyl, aryl and heteroaryl, each of which is optionally substituted with one, two or three groups independently selected from lower alkyl, halogen, lower alkoxy, hydroxy, amino, mono- or dialkylamino, and trifluoromethyl;

and pharmaceutically acceptable derivatives thereof.

29. The compound of Claim 28 wherein R' is selected from aminomethyl, aminoethyl, aminopropyl, isopropylaminomethyl, t-butylaminomethyl, iso-butylaminomethyl, 1-methylpropylaminomethyl, 2-

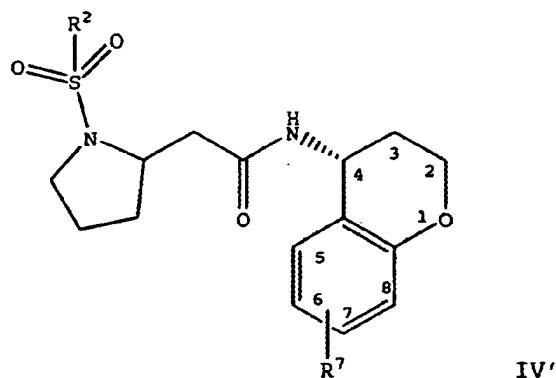
methylbutylaminomethyl, 2,2'-dimethylpropylaminomethyl,
2,2',3-trimethylpropylaminomethyl, allyl-aminomethyl,
isopropylaminopropyl, 1-(isobutylamino)ethyl, 1-
(isopropylamino)-1-methylethyl, N-isopropyl-N-
5 ethylaminomethyl, N-isopropyl-N-methylaminomethyl, N-t-
butyl-N-methylaminomethyl, N-iso-butyl-N-methylaminomethyl,
N-t-butyl-N-ethylaminomethyl, N-isobutyl-N-
methylaminomethyl, N-t-butyl-N-isopropylaminomethyl, N,N-
di(isopropyl)aminomethyl, N,N-dimethylaminomethyl, N,N-
10 diethylaminomethyl, N,N-di(t-butyl)-aminomethyl, N,N-
di(allyl)-aminomethyl, cyclopropylaminomethyl, 1-
(cyclopropylamino)ethyl, cyclobutylaminomethyl, 2-
(cyclobutylamino)ethyl, 1-(cyclobutylamino)ethyl,
cyclopentylaminomethyl, 1-cyclopentylaminoethyl,
15 cyclopropylmethylaminomethyl, hydroxyethylamino-allyl,
isopropylamino-allyl, t-butylamino-allyl,
cyclopropylmethylamino-allyl, piperidin-1-yl-allyl,
pyrrolidin-1-yl-allyl, azetidin-1-yl-allyl, 3-
hydroxypyrrolidin-1-yl-allyl, aminocarbonylethylaminomethyl,
20 methoxyethylaminomethyl, 1-(methoxyethylamino)ethyl, 1-
piperidinylmethyl, 2-(piperidin-1-yl)ethyl, 3,4-
dihdropiperidin-1-ylmethyl, 4-fluoropiperidinylmethyl,
4,4'-difluoropiperidinylmethyl, 4-(piperidin-1-
yl)piperidinylmethyl, 3-aminocarbonylpiperidin-1-ylmethyl,
25 4-(dimethylamino)piperidin-1-ylmethyl, 2,6-
dimethylpiperidin-1-ylmethyl, 3,3-dimethylpiperidin-1-
ylmethyl, piperidin-1-yl-2-methylethyl, 3-hydroxypiperidin-
1-yl, 4-morpholinylmethyl, 4-morpholinylethyl, 1-
pyrrolidinylmethyl, 2-methylpyrrolidin-1-ylmethyl, 1-
30 (methylpyrrolidin-1-yl)ethyl, 2,5-dimethylpyrrolidin-1-
ylmethyl, 1-azetidinylmethyl, 7-aza-bicyclo[2.2.1]heptyl,
piperazin-1-ylmethyl, 4-methylpiperazin-1-ylmethyl, and 1-
pyrrolidinylethylaminomethyl;
and pharmaceutically acceptable derivatives thereof.

30. The compound of Claim 28 wherein R⁷ is at position 7.

5 31. The compound of Claim 28 wherein R² is 2-naphthyl, 3,4-dichlorophenyl or 3-trifluoromethylphenyl.

32. A compound of Formula IV'

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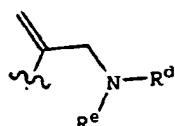


IV'

wherein R² is selected from naphthyl, phenyl, pyridinyl, benzothienyl, quinolinyl and isoquinolinyl, and wherein each is optionally substituted with one to three substituents selected from chloro, fluoro, methoxy, methyl, trifluoromethyl, and phenyl; and

15

20

wherein R⁷ is selected from  , C₃₋₆-cycloalkyl (C₁-C₂)alkylamino (C₁-C₂)alkyl, C₃₋₆-cycloalkylamino (C₁-C₂)alkyl, (C₁-C₂)alkoxy (C₁-C₂)alkylamino (C₁-C₂)alkyl, mono-C₂₋₄-alkenylamino-C₁₋₄-alkyl, di-C₂₋₄-alkenylamino-C₁₋₄-alkyl, hydroxy-C₁₋₄-alkylamino-C₁₋₄-alkyl, aminocarbonyl-C₁₋₄-alkylamino-C₁₋₂-alkyl, mono-C₁₋₆-alkylamino-C₁₋₄-alkyl, di-C₁₋₄-alkylamino-C₁₋₄-alkyl and 5-8 membered heterocyclyl-C₁₋₄-alkyl; wherein the 5-8 membered heterocyclyl-(CH₂)_p-

optionally substituted with one to three groups independently selected from halo, -NH₂, -OH, -CN, -CF₃, (C₁-C₆)alkylamino, oxo, (C₁-C₆)alkoxy, (C₂-C₆)alkenyl, (C₂-C₆)alkynyl, di(C₁-C₆)alkylamino, -C(O)R⁸, -COOR⁸, -C(O)NR⁸R^{8'}, -NR⁸C(O)R^{8'}, =NCN;

5 wherein R⁴ is selected from C₁₋₅-alkyl, C₃₋₆-cycloalkyl, C₃₋₆-cycloalkyl-C₁₋₃-alkyl, C₁₋₄-hydroxyalkyl, C₁₋₃-alkoxy-C₁₋₃-alkyl and H; and

wherein R⁶ is H; or where R⁴ and R⁶ together with the

10 nitrogen atom to which they are attached form a 4-8 membered nitrogen-containing heterocyclic ring;

wherein R⁷ is at position 6, 7 or 8; and

wherein R⁸ and R^{8'} independently are selected from H, and lower alkyl, aryl and heteroaryl, each of which is

15 optionally substituted with one, two or three groups independently selected from lower alkyl, halogen, lower alkoxy, hydroxy, amino, mono- or dialkylamino, and trifluoromethyl;

and pharmaceutically acceptable derivatives thereof.

20 33. The compound of Claim 32 wherein R⁷ is selected from aminomethyl, aminoethyl, aminopropyl, isopropylaminomethyl, t-butylaminomethyl, iso-butylaminomethyl, 1-methylpropylaminomethyl, 2-methylbutylaminomethyl, 2,2'-dimethylpropylaminomethyl, 25 2,2',3-trimethylpropylaminomethyl, allyl-aminomethyl, isopropylaminopropyl, 1-(isobutylamino)ethyl, 1-(isopropylamino)-1-methylethyl, N-isopropyl-N-ethylaminomethyl, N-isopropyl-N-methylaminomethyl, N-t-butyl-N-methylaminomethyl, N-iso-butyl-N-methylaminomethyl, 30 N-t-butyl-N-ethylaminomethyl, N-isobutyl-N-methylaminomethyl, N-t-butyl-N-isopropylaminomethyl, N,N-di(isopropyl)aminomethyl, N,N-dimethylaminomethyl, N,N-diethylaminomethyl, N,N-di(t-butyl)-aminomethyl, N,N-

di(allyl)-aminomethyl, cyclopropylaminomethyl, 1-(cyclopropylamino)ethyl, cyclobutylaminomethyl, 2-(cyclobutylamino)ethyl, 1-(cyclobutylamino)ethyl, cyclopentylaminomethyl, 1-cyclopentylaminoethyl,

5 cyclopropylmethylaminomethyl, hydroxyethylamino-allyl, isopropylamino-allyl, t-butylamino-allyl, cyclopropylmethylamino-allyl, piperidin-1-yl-allyl, pyrrolidin-1-yl-allyl, azetidin-1-yl-allyl, 3-hydroxypyrrolidin-1-yl-allyl, aminocarbonylethylaminomethyl,

10 methoxyethylaminomethyl, 1-(methoxyethylamino)ethyl, 1-piperidinylmethyl, 2-(piperidin-1-yl)ethyl, 3,4-dihydropiperidin-1-ylmethyl, 4-fluoropiperidinylmethyl, 4,4'-difluoropiperidinylmethyl, 4-(piperidin-1-yl)piperidinylmethyl, 3-aminocarbonylpiperidin-1-ylmethyl,

15 4-(dimethylamino)piperidin-1-ylmethyl, 2,6-dimethylpiperidin-1-ylmethyl, 3,3-dimethylpiperidin-1-ylmethyl, piperidin-1-yl-2-methylethyl, 3-hydroxypiperidin-1-yl, 4-morpholinylmethyl, 4-morpholinylethyl, 1-pyrrolidinylmethyl, 2-methylpyrrolidin-1-ylmethyl, 1-(methylpyrrolidin-1-yl)ethyl, 2,5-dimethylpyrrolidin-1-ylmethyl, 1-azetidinylmethyl, 7-aza-bicyclo[2.2.1]heptyl, piperazin-1-ylmethyl, 4-methylpiperazin-1-ylmethyl, and 1-pyrrolidinylethylaminomethyl;

20 and pharmaceutically acceptable derivatives thereof.

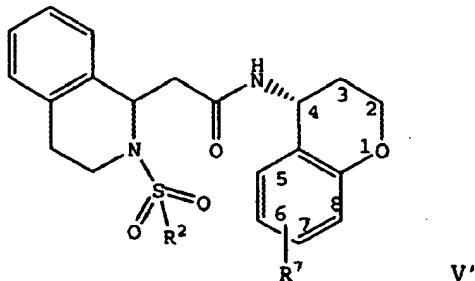
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34. The compound of Claim 32 wherein R is at position 7.

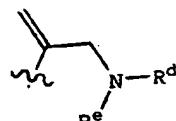
35. The compound of Claim 32 wherein R² is 2-naphthyl,

30 3,4-dichlorophenyl or 3-trifluoromethylphenyl.

36. A compound of Formula V'



5 wherein R^2 is selected from naphthyl, phenyl, pyridinyl, benzothienyl, quinolinyl and isoquinolinyl, and wherein each is optionally substituted with one to three substituents selected from chloro, fluoro, methoxy, methyl, trifluoromethyl, and phenyl; and



10 wherein R^7 is selected from R^e , C_{3-6} -cycloalkyl (C_1-C_2) alkylamino (C_1-C_2) alkyl, C_{3-6} -cycloalkylamino (C_1-C_2) alkyl, (C_1-C_2) alkoxy (C_1-C_2) alkylamino (C_1-C_2) alkyl, mono- C_{2-4} -alkenylamino- C_{1-4} -alkyl, di- C_{2-4} -alkenylamino- C_{1-4} -alkyl, hydroxy- C_{1-4} -alkylamino- C_{1-4} -alkyl, aminocarbonyl- C_{1-4} -alkylamino- C_{1-2} -alkyl, mono- C_{1-6} -alkylamino- C_{1-4} -alkyl, di- C_{1-4} -alkylamino- C_{1-4} -alkyl and 5-8 membered heterocyclyl- C_{1-4} -alkyl; wherein the 5-8 membered heterocyclyl- $(CH_2)_p$ - is optionally substituted with one to three groups independently selected from halo, -NH₂, -OH, -CN, -CF₃, (C_1-C_6) alkylamino, oxo, (C_1-C_6) alkoxy, (C_2-C_6) alkenyl, (C_2-C_6) alkynyl, di(C_1-C_6) alkylamino, -C(O)R⁸, -COOR⁸, -C(O)NR⁸R⁸, -NR⁸C(O)R⁸, =NCR⁸; wherein R⁴ is selected from C_{1-5} -alkyl, C_{3-6} -cycloalkyl, C_{3-6} -cycloalkyl- C_{1-4} -alkyl, C_{1-4} -hydroxyalkyl, C_{1-3} -alkoxy- C_{1-3} -alkyl and H; and

wherein R^e is H; or where R^d and R^e together with the nitrogen atom to which they are attached form a 4-8 membered nitrogen-containing heterocyclic ring;

wherein R⁷ is at position 6, 7 or 8; and

5 wherein R⁸ and R^{8'} independently are selected from H, and lower alkyl, aryl and heteroaryl, each of which is optionally substituted with one, two or three groups independently selected from lower alkyl, halogen, lower alkoxy, hydroxy, amino, mono- or dialkylamino, 10 and trifluoromethyl;

and pharmaceutically acceptable derivatives thereof.

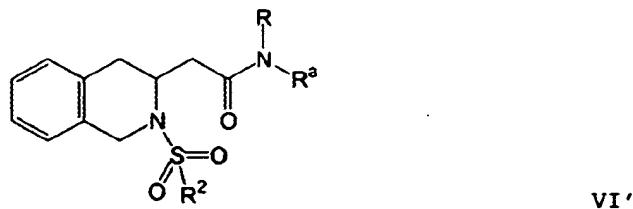
37. The compound of Claim 36 wherein R⁷ is selected from aminomethyl, aminoethyl, aminopropyl, 15 isopropylaminomethyl, t-butylaminomethyl, iso-butylaminomethyl, 1-methylpropylaminomethyl, 2-methylbutylaminomethyl, 2,2'-dimethylpropylaminomethyl, 2,2',3-trimethylpropylaminomethyl, allyl-aminomethyl, isopropylaminopropyl, 1-(isobutylamino)ethyl, 1-20 (isopropylamino)-1-methylethyl, N-isopropyl-N-ethylaminomethyl, N-isopropyl-N-methylaminomethyl, N-t-butyl-N-methylaminomethyl, N-iso-butyl-N-methylaminomethyl, N-t-butyl-N-ethylaminomethyl, N-isobutyl-N-methylaminomethyl, N-t-butyl-N-isopropylaminomethyl, N,N-25 di(isopropyl)aminomethyl, N,N-dimethylaminomethyl, N,N-diethylaminomethyl, N,N-di(t-butyl)-aminomethyl, N,N-di(allyl)-aminomethyl, cyclopropylaminomethyl, 1-(cyclopropylamino)ethyl, cyclobutylaminomethyl, 2-(cyclobutylamino)ethyl, 1-(cyclobutylamino)ethyl, 30 cyclopentylaminomethyl, 1-cyclopentylaminoethyl, cyclopropylmethylaminomethyl, hydroxyethylamino-allyl, isopropylamino-allyl, t-butylamino-allyl, cyclopropylmethylamino-allyl, piperidin-1-yl-allyl, pyrrolidin-1-yl-allyl, azetidin-1-yl-allyl, 3-

hydroxypyrrolidin-1-yl-allyl, aminocarbonylethylaminomethyl, methoxyethylaminomethyl, 1-(methoxyethylamino)ethyl, 1-piperidinylmethyl, 2-(piperidin-1-yl)ethyl, 3,4-dihdropiperidin-1-ylmethyl, 4-fluoropiperidinylmethyl,
 5 4,4'-difluoropiperidinylmethyl, 4-(piperidin-1-yl)piperidinylmethyl, 3-aminocarbonylpiperidin-1-ylmethyl, 4-(dimethylamino)piperidin-1-ylmethyl, 2,6-dimethylpiperidin-1-ylmethyl, 3,3-dimethylpiperidin-1-ylmethyl, piperidin-1-yl-2-methylethyl, 3-hydroxypiperidin-10-yl, 4-morpholinylmethyl, 4-morpholinylethyl, 1-pyrrolidinylmethyl, 2-methylpyrrolidin-1-ylmethyl, 1-(methylpyrrolidin-1-yl)ethyl, 2,5-dimethylpyrrolidin-1-ylmethyl, 1-azetidinylmethyl, 7-aza-bicyclo[2.2.1]heptyl, piperazin-1-ylmethyl, 4-methylpiperazin-1-ylmethyl, and 1-pyrrolidinylethylaminomethyl;
 15 and pharmaceutically acceptable derivatives thereof.

38. The compound of Claim 36 wherein R is at position 7.
 20

39. The compound of Claim 36 wherein R² is 2-naphthyl, 3,4-dichlorophenyl or 3-trifluoromethylphenyl.

40. A compound of Formula VI'
 25



wherein R is a 9-11 membered fused bicyclic carbocyclic or heterocyclic ring substituted with one to three basic 30 moieties, and optionally substituted with one to three

groups independently selected from halo, -NH₂, -OH, -CN, -CF₃, (C₁-C₆)alkylamino, oxo, (C₁-C₆)alkoxy, (C₂-C₆)alkenyl, (C₂-C₆)alkynyl, di(C₁-C₆)alkylamino, -C(O)R⁸, -COOR⁸, -C(O)NR⁸R^{8'}, -NR⁸C(O)R^{8'}, and

5 (C₁-C₆)alkyl, aryl, heteroaryl, cycloalkyl or heterocyclyl, each of which is optionally substituted with one to three groups independently selected from halo, -NH₂, -OH, -CN, -CF₃, (C₁-C₆)alkylamino, halo(C₁-C₆)alkyl, oxo, (C₁-C₆)alkoxy, (C₁-C₆)alkoxy(C₁-C₆)alkyl, (C₁-C₆)alkyl, (C₂-C₆)alkenyl, (C₂-C₆)alkynyl, di(C₁-C₆)alkylamino, -C(O)R⁸, -COOR⁸, -C(O)NR⁸R^{8'}, and -NR⁸C(O)R^{8'};

10 wherein R⁸ and R^{8'} independently are selected from H, and lower alkyl, aryl and heteroaryl, each of which is optionally substituted with one, two or three groups independently selected from lower alkyl, halogen, lower alkoxy, hydroxy, amino, mono- or dialkylamino, and trifluoromethyl;

15 wherein R² is selected from arylalkenyl, aryl, and heterocyclyl, wherein R² is optionally substituted with one to five groups independently selected from halo, -NH₂, -OH, -CN, -CF₃, (C₁-C₆)alkylamino, oxo, (C₁-C₆)alkoxy, (C₂-C₆)alkenyl, (C₂-C₆)alkynyl, di(C₁-C₆)alkylamino, -C(O)R⁸, -COOR⁸, -C(O)NR⁸R^{8'}, -NR⁸C(O)R^{8'}, and

20 (C₁-C₆)alkyl, aryl, heteroaryl, cycloalkyl and heterocyclyl, each of which is optionally substituted with one to three groups independently selected from halo, -NH₂, -OH, -CN, -CF₃, (C₁-C₆)alkylamino, halo(C₁-C₆)alkyl, oxo, (C₁-C₆)alkoxy, (C₁-C₆)alkoxy(C₁-C₆)alkyl, (C₁-C₆)alkyl, (C₂-C₆)alkenyl, (C₂-C₆)alkynyl, di(C₁-C₆)alkylamino, -C(O)R⁸, -COOR⁸, -C(O)NR⁸R^{8'}, and -NR⁸C(O)R^{8'};

25 and (C₁-C₆)alkyl, aryl, heteroaryl, cycloalkyl and heterocyclyl, each of which is optionally substituted with one to three groups independently selected from halo, -NH₂, -OH, -CN, -CF₃, (C₁-C₆)alkylamino, halo(C₁-C₆)alkyl, oxo, (C₁-C₆)alkoxy, (C₁-C₆)alkoxy(C₁-C₆)alkyl, (C₁-C₆)alkyl, (C₂-C₆)alkenyl, (C₂-C₆)alkynyl, di(C₁-C₆)alkylamino, -C(O)R⁸, -COOR⁸, -C(O)NR⁸R^{8'}, and -NR⁸C(O)R^{8'};

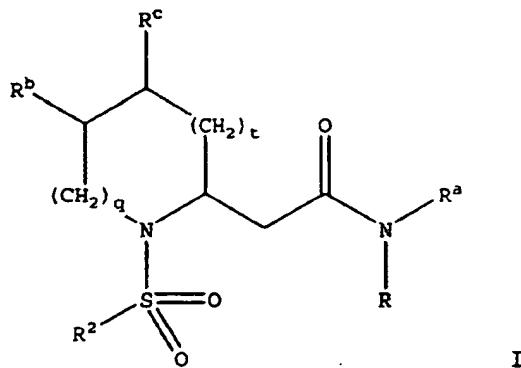
30 and (C₁-C₆)alkyl, aryl, heteroaryl, cycloalkyl and heterocyclyl, each of which is optionally substituted with one to three groups independently selected from halo, -NH₂, -OH, -CN, -CF₃, (C₁-C₆)alkylamino, halo(C₁-C₆)alkyl, oxo, (C₁-C₆)alkoxy, (C₁-C₆)alkoxy(C₁-C₆)alkyl, (C₁-C₆)alkyl, (C₂-C₆)alkenyl, (C₂-C₆)alkynyl, di(C₁-C₆)alkylamino, -C(O)R⁸, -COOR⁸, -C(O)NR⁸R^{8'}, and -NR⁸C(O)R^{8'};

wherein R^a is independently selected from H and C₁₋₄-alkyl,
 and aryl optionally substituted with one to three groups
 independently selected from halo, -NH₂, -OH, -CN, -CF₃,
 (C₁-C₆)alkylamino, halo(C₁-C₆)alkyl, oxo, (C₁-C₆)alkoxy,
 5 (C₁-C₆)alkoxy(C₁-C₆)alkyl, (C₁-C₆)alkyl, (C₂-C₆)alkenyl,
 (C₂-C₆)alkynyl, di(C₁-C₆)alkylamino, -C(O)R^b, -COOR^b,
 -C(O)NR^bR^b, and -NR^bC(O)R^b.

41. The compound of Claim 40 wherein R is selected
 10 from 1,2,3,4-tetrahydronaphth-1-yl, 1,2,3,4-
 tetrahydronaphth-2-yl, indan-1-yl and indan-2-yl, chroman-4-
 yl, and 2,2-dioxo-3,4-dihydro-1H-2, 1-benzothiazin-4-yl.

42. The compound of Claim 40 R^a is selected from H
 15 and (C₁-C₂)alkyl.

43. A compound of Formula I



20 wherein q is 0-3;
 wherein t is 0-2, provided that when t is 2, q is not 3;
 wherein R is a 9-11 membered fused carbocyclic or
 heterocyclic ring substituted with one to three basic
 25 moieties, and optionally substituted with one to three
 groups independently selected from halo, -NH₂, -OH, -CN,
 -CF₃, (C₁-C₆)alkylamino, oxo, (C₁-C₆)alkoxy, (C₂-

C_6) alkenyl, (C_2-C_6) alkynyl, di(C_1-C_6) alkylamino, $-C(O)R^8$, $-COOR^8$, $-C(O)NR^8R^8'$, $-NR^8C(O)R^8'$, and (C_1-C_6) alkyl, aryl, heteroaryl, cycloalkyl or heterocyclyl, each of which is optionally substituted with one to three groups independently selected from halo, $-NH_2$, $-OH$, $-CN$, $-CF_3$, (C_1-C_6) alkylamino, halo(C_1-C_6) alkyl, oxo, (C_1-C_6) alkoxy, (C_1-C_6) alkoxy(C_1-C_6) alkyl, (C_1-C_6) alkyl, (C_2-C_6) alkenyl, (C_2-C_6) alkynyl, di(C_1-C_6) alkylamino, $-C(O)R^8$, $-COOR^8$, $-C(O)NR^8R^8'$, and $-NR^8C(O)R^8'$;

wherein R^8 and R^8' independently are selected from H, and lower alkyl, aryl and heteroaryl, each of which is optionally substituted with one, two or three groups independently selected from lower alkyl, halogen, lower alkoxy, hydroxy, amino, mono- or dialkylamino, and trifluoromethyl;

wherein R^2 is selected from arylalkenyl, aryl, and heterocyclyl, wherein R^2 is optionally substituted with one to five groups independently selected from halo, $-NH_2$, $-OH$, $-CN$, $-CF_3$, (C_1-C_6) alkylamino, oxo, (C_1-C_6) alkoxy, (C_2-C_6) alkenyl, (C_2-C_6) alkynyl, di(C_1-C_6) alkylamino, $-C(O)R^8$, $-COOR^8$, $-C(O)NR^8R^8'$, $-NR^8C(O)R^8'$, and (C_1-C_6) alkyl, aryl, heteroaryl, cycloalkyl and heterocyclyl, each of which is optionally substituted with one to three groups independently selected from halo, $-NH_2$, $-OH$, $-CN$, $-CF_3$, (C_1-C_6) alkylamino, halo(C_1-C_6) alkyl, oxo, (C_1-C_6) alkoxy, (C_1-C_6) alkoxy(C_1-C_6) alkyl, (C_1-C_6) alkyl, (C_2-C_6) alkenyl, (C_2-C_6) alkynyl, di(C_1-C_6) alkylamino, $-C(O)R^8$, $-COOR^8$, $-C(O)NR^8R^8'$, and $-NR^8C(O)R^8'$;

wherein R^a is independently selected from H and C_{1-4} -alkyl, and

aryl optionally substituted with one to three groups independently selected from halo, -NH₂, -OH, -CN, -CF₃, (C₁-C₆)alkylamino, halo(C₁-C₆)alkyl, oxo, (C₁-C₆)alkoxy, (C₁-C₆)alkoxy(C₁-C₆)alkyl, (C₁-C₆)alkyl, (C₂-C₆)alkenyl, (C₂-C₆)alkynyl, di(C₁-C₆)alkylamino, -C(O)R^a, -COOR^a, -C(O)NR^aR^{a'}, and -NR^aC(O)R^{a'};

5 wherein R^b is independently selected from H and C₁₋₂-alkyl; and

wherein R^c is independently selected from H and C₁₋₂-alkyl;

10 or

wherein R^b and R^c may be joined to form a 6-membered aryl or heteroaryl ring optionally substituted with one to three groups independently selected from halo, -NH₂, -OH, -CN, -CF₃, (C₁-C₆)alkylamino, oxo, (C₁-C₆)alkoxy, (C₁-C₆)alkenyl, (C₂-C₆)alkynyl, di(C₁-C₆)alkylamino, -C(O)R^a, -COOR^a, -C(O)NR^aR^{a'}, -NR^aC(O)R^{a'}, and

15 (C₁-C₆)alkyl, aryl, heteroaryl, cycloalkyl and heterocyclyl, each of which is optionally substituted with one to three groups independently selected from halo, -NH₂, -OH, -CN, -CF₃, (C₁-C₆)alkylamino, halo(C₁-C₆)alkyl, oxo, (C₁-C₆)alkoxy, (C₁-C₆)alkoxy(C₁-C₆)alkyl, (C₁-C₆)alkyl, (C₂-C₆)alkenyl, (C₂-C₆)alkynyl, di(C₁-C₆)alkylamino, -C(O)R^a, -COOR^a, -C(O)NR^aR^{a'}, and -NR^aC(O)R^{a'};

20 25 and pharmaceutically acceptable derivatives thereof; provided the basic moiety is not 2-oxo-piperaziny-4-ylmethyl.

44. The compound of Claim 1 and/or pharmaceutically acceptable derivatives thereof selected from

N-(7-Piperidin-1-ylmethyl-chroman-4-(R)-yl)-2-[1-(3-trifluoromethyl-benzenesulfonyl)-piperidin-2-yl]-acetamide;

2-[(1-(Naphthalene-2-sulfonyl)-piperidin-2-yl)-N-(7-piperidin-1-ylmethyl-chroman-4-(R)-yl)-acetamide;

2-[(1-(Naphthalene-2-sulfonyl)-pyrrolidin-2-(L)-yl)-N-(7-piperidin-1-ylmethyl-chroman-4-(R)-yl)-acetamide;

5 N-((1R)-6-(1-piperidinylmethyl)-1,2,3,4-tetrahydro-1-naphthalenyl)-2-((2S)-1-((3-(trifluoromethyl)phenyl)sulfonyl)-2;

N-((1R)-6-(((1,1-dimethylethyl)amino)methyl)-1,2,3,4-tetrahydro-1-naphthalenyl)-2-((2S)-1-((3-(trifluoromethyl)phenyl)sulfonyl)-2-

10 piperidinyl)acetamide;

N-((1R)-6-(((1,1-dimethylethyl)amino)methyl)-1,2,3,4-tetrahydro-1-naphthalenyl)-2-((2S)-1-((4-methylphenyl)sulfonyl)-2-piperidinyl)acetamide;

15 2-((2S)-1-((3-chloro-4-methylphenyl)sulfonyl)-2-piperidinyl)-N-((1R)-6-(((1,1-dimethylethyl)amino)methyl)-1,2,3,4-tetrahydro-1-naphthalenyl)acetamide;

2-((2S)-1-((2,4,6-trimethylphenyl)sulfonyl)-2-piperidinyl)-

20 N-((1R)-6-(((1,1-dimethylethyl)amino)methyl)-1,2,3,4-tetrahydro-1-naphthalenyl)acetamide;

N-((1R)-6-(1-piperidinylmethyl)-1,2,3,4-tetrahydro-1-naphthalenyl)-2-((2S)-1-((2,4,6-trimethylphenyl)sulfonyl)-1,2,3,4-tetrahydro-2-

25 quinolinyl)acetamide;

2-((2S)-1-((3,4-dichlorophenyl)sulfonyl)-2-piperidinyl)-N-((1R)-6-(((1,1-dimethylethyl)amino)methyl)-1,2,3,4-tetrahydro-1-naphthalenyl)acetamide;

N-((1R)-6-(1-piperidinyl)methyl)-1,2,3,4-tetrahydro-1-

30 naphthalenyl)-2-((2S)-1-((3-(trifluoromethyl)phenyl)sulfonyl)-2-piperidinyl)acetamide;

N-((1R)-6-((cyclobutylamino)methyl)-1,2,3,4-tetrahydro-1-naphthalenyl)-2-((2S)-1-((3-

(trifluoromethyl)phenyl)sulfonyl)-2-
piperidinyl)acetamide;

N-methyl-N-((4R)-7-(1-piperidinylmethyl)-3,4-dihydro-2H-
chromen-4-yl)-2-((2S)-1-((3-
5 (trifluoromethyl)phenyl)sulfonyl)-2-
piperidinyl)acetamide;

N-((1R)-6-(((1,1-dimethylethyl)amino)methyl)-1,2,3,4-
tetrahydro-1-naphthalenyl)-2-((2S)-1-((4-(1,1-
dimethylethyl)phenyl)sulfonyl)-2-piperidinyl)acetamide;

10 N-((1R)-6-(((1,1-dimethylethyl)amino)methyl)-1,2,3,4-
tetrahydro-1-naphthalenyl)-2-((2S)-1-((4-(1,1-
dimethylethyl)phenyl)sulfonyl)-2-piperidinyl)acetamide;

N-((1R)-6-((diethylamino)methyl)-1,2,3,4-tetrahydro-1-
naphthalenyl)-2-((2S)-1-((3-
15 (trifluoromethyl)phenyl)sulfonyl)-2-
piperidinyl)acetamide;

N-((1R)-6-((isobutylamino)methyl)-1,2,3,4-tetrahydro-1-
naphthalenyl)-2-((2S)-1-((3-
(trifluoromethyl)phenyl)sulfonyl)-2-
20 piperidinyl)acetamide;

N-((1R)-6-(((1,1-dimethylethyl)amino)methyl)-1,2,3,4-
tetrahydro-1-naphthalenyl)-2-((2S)-1-((4-methyl-3-
(trifluoromethyl)phenyl)sulfonyl)-2-
piperidinyl)acetamide;

25 N-((1R)-6-((cyclopropylamino)methyl)-1,2,3,4-tetrahydro-1-
naphthalenyl)-2-((2S)-1-((3-
(trifluoromethyl)phenyl)sulfonyl)-2-
piperidinyl)acetamide;

N-((1R)-6-(((2-methylbutyl)amino)methyl)-1,2,3,4-tetrahydro-
30 1-naphthalenyl)-2-((2S)-1-((3-
(trifluoromethyl)phenyl)sulfonyl)-2-
piperidinyl)acetamide;

N-((1R)-6-(((2-(methyloxy)ethyl)amino)methyl)-1,2,3,4-
tetrahydro-1-naphthalenyl)-2-((2S)-1-((3-

(trifluoromethyl)phenyl)sulfonyl)-2-
piperidinyl)acetamide;

N-((1R)-6-(((cyclopropylmethyl)amino)methyl)-1,2,3,4-
tetrahydro-1-naphthalenyl)-2-((2S)-1-((3-
5 (trifluoromethyl)phenyl)sulfonyl)-2-
piperidinyl)acetamide;

N-((1R)-6-(((isopropylmethyl)amino)methyl)-1,2,3,4-
tetrahydro-1-naphthalenyl)-2-((2S)-1-((3-
10 (trifluoromethyl)phenyl)sulfonyl)-2-
piperidinyl)acetamide;

N-((1R)-6-((4-fluoro-1-piperidinyl)methyl)-1,2,3,4-
tetrahydro-1-naphthalenyl)-2-((2S)-1-((3-
15 (trifluoromethyl)phenyl)sulfonyl)-2-
piperidinyl)acetamide;

N-((1R)-6-(((1,1-dimethylethyl)amino)methyl)-1,2,3,4-
tetrahydro-1-naphthalenyl)-2-((2R/S)-1-((3-
20 (trifluoromethyl)phenyl)sulfonyl)-1,2,3,4-tetrahydro-2-
quinolinyl)acetamide;

N-((1R)-6-((4-fluoro-1-piperidinyl)methyl)-1,2,3,4-
25 tetrahydro-1-naphthalenyl)-2-((2R/S)-1-((3-
(trifluoromethyl)phenyl)sulfonyl)-1,2,3,4-tetrahydro-2-
quinolinyl)acetamide;

N-((1R)-6-(((cyclopropylmethyl)amino)methyl)-1,2,3,4-
tetrahydro-1-naphthalenyl)-2-((2R/S)-1-((3-
30 (trifluoromethyl)phenyl)sulfonyl)-1,2,3,4-tetrahydro-2-
quinolinyl)acetamide;

N-((1R)-6-(((isobutylmethyl)amino)methyl)-1,2,3,4-
tetrahydro-1-naphthalenyl)-2-((2R/S)-1-((3-
(trifluoromethyl)phenyl)sulfonyl)-1,2,3,4-tetrahydro-2-
quinolinyl)acetamide;

N-((1R)-6-(((diethylamino)methyl)-1,2,3,4-tetrahydro-1-naphthalenyl)-2-((2R/S)-1-((3-(trifluoromethyl)phenyl)sulfonyl)-1,2,3,4-tetrahydro-2-quinolinyl)acetamide;

5 N-((1R)-6-((4-fluoro-1-piperidinyl)methyl)-1,2,3,4-tetrahydro-1-naphthalenyl)-2-((2R)-1-((4-methylphenyl)sulfonyl)-1,2,3,4-tetrahydro-2-quinolinyl)acetamide;

10 2-((2R/S)-1-((4-methylphenyl)sulfonyl)-1,2,3,4-tetrahydro-2-quinolinyl)-N-((1R)-6-(((2-methylpropyl)amino)methyl)-1,2,3,4-tetrahydro-1-naphthalenyl)acetamide;

15 N-((1R)-6-(((2,2-dimethylpropyl)amino)methyl)-1,2,3,4-tetrahydro-1-naphthalenyl)-2-((2S)-1-((3-(trifluoromethyl)phenyl)sulfonyl)-2-piperidinyl)acetamide;

20 2-((2S)-1-(1-benzothien-3-ylsulfonyl)-2-piperidinyl)-N-((1R)-6-(((1,1-dimethylethyl)amino)methyl)-1,2,3,4-tetrahydro-1-naphthalenyl)acetamide;

25 2-((2S)-1-(1-benzothien-3-ylsulfonyl)-2-piperidinyl)-N-((1R)-6-(((2,2-dimethylpropyl)amino)methyl)-1,2,3,4-tetrahydro-1-naphthalenyl)acetamide;

30 1-(((5R)-5-(((2S)-1-((3-(trifluoromethyl)phenyl)sulfonyl)-2-piperidinyl)acetyl)amino)-5,6,7,8-tetrahydro-2-naphthalenyl)methyl)-3-piperidinecarboxamide;

35 N-((4R)-7-(4-morpholinylmethyl)-3,4-dihydro-2H-chromen-4-yl)-2-((1S)-2-((3-(trifluoromethyl)phenyl)sulfonyl)-1,2,3,4-tetrahydro-1-isoquinolinyl)acetamide;

40 N-((4R)-7-(7-azabicyclo[2.2.1]hept-7-ylmethyl)-3,4-dihydro-2H-chromen-4-yl)-2-((1S)-2-((3-(trifluoromethyl)phenyl)sulfonyl)-1,2,3,4-tetrahydro-1-isoquinolinyl)acetamide;

45 N-((4R)-7-(1-Piperidinylmethyl)-3,4-dihydro-2H-chromen-4-yl)-2-((1R)-2-((3-(trifluoromethyl)phenyl)sulfonyl)-1,2,3,4-tetrahydro-1-isoquinolinyl)acetamide;

N-((4R)-7-((4-Fluoro-1-piperidinyl)methyl)-3,4-dihydro-2H-chromen-4-yl)-2-((2S)-1-((3-(trifluoromethyl)phenyl)sulfonyl)-2-pyrrolidinyl)acetamide;

N-((4R)-7-((4,4-Difluoro-1-piperidinyl)methyl)-3,4-dihydro-2H-chromen-4-yl)-2-((2S)-1-((3-(trifluoromethyl)phenyl)sulfonyl)-2-pyrrolidinyl)acetamide;

5 N-((4R)-7-((2-Naphthalenylsulfonyl)-2-piperidinyl)-N-((4R)-7-(1-piperidinylmethyl)-3,4-dihydro-2H-chromen-4-yl)acetamide;

10 N-((4R)-6-chloro-7-(1-piperidinylmethyl)-3,4-dihydro-2H-chromen-4-yl)-2-((2S)-1-((3-(trifluoromethyl)phenyl)sulfonyl)-2-pyrrolidinyl)acetamide;

N-((4R)-7-(1-Piperidinylmethyl)-3,4-dihydro-2H-chromen-4-yl)-2-((3R)-2-((3-(trifluoromethyl)phenyl)sulfonyl)-1,2,3,4-tetrahydro-3-isoquinolinyl)acetamide;

15 N-((4R)-7-(1-Piperidinylmethyl)-3,4-dihydro-2H-chromen-4-yl)-2-((1S)-2-((3-(trifluoromethyl)phenyl)sulfonyl)-1,2,3,4-tetrahydro-1-isoquinolinyl)acetamide;

N-((4R)-7-(4-Morpholinylmethyl)-3,4-dihydro-2H-chromen-4-yl)-2-((1S)-2-((3-(trifluoromethyl)phenyl)sulfonyl)-1,2,3,4-tetrahydro-1-isoquinolinyl)acetamide;

20 N-((4R)-7-(7-Azabicyclo[2.2.1]hept-7-ylmethyl)-3,4-dihydro-2H-chromen-4-yl)-2-((1S)-2-((3-(trifluoromethyl)phenyl)sulfonyl)-1,2,3,4-tetrahydro-1-isoquinolinyl)acetamide;

N-((4R)-7-(1-Piperidinylmethyl)-3,4-dihydro-2H-chromen-4-yl)-2-((2S)-1-((3-(trifluoromethyl)phenyl)sulfonyl)-2-piperidinyl)acetamide;

25 N-((4R)-7-(1-Piperidinylmethyl)-3,4-dihydro-2H-chromen-4-yl)-2-((2R)-1-((3-(trifluoromethyl)phenyl)sulfonyl)-2-piperidinyl)acetamide;

N-((1R)-6-((1S)-1-methyl-2-(1-piperidinyl)ethyl)-1,2,3,4-tetrahydro-1-naphthalenyl)-2-((2S)-1-((3-

(trifluoromethyl)phenyl)sulfonyl)-2-
piperidinyl)acetamide; and
N-((1R)-6-(1-(1-piperidinylmethyl)ethenyl)-1,2,3,4-
tetrahydro-1-naphthalenyl)-2-((2S)-1-((3-
5 (trifluoromethyl)phenyl)sulfonyl)-2-
piperidinyl)acetamide.

45. A pharmaceutically acceptable salt of a compound
of Claim 1.

10

46. A pharmaceutical composition comprising a
pharmaceutically acceptable carrier and a compound of Claim
1.

15 47.

46. A method of treating pain comprising administering
an effective amount of a compound of Claim 1.

20

48. 47. A pharmaceutical composition for the treatment of
disease conditions mediated by bradykinin, in a mammalian
subject, which comprises a therapeutically effective amount
of a compound according to Claim 1 and a pharmaceutically
acceptable carrier.

25

49. 48. A pharmaceutical composition for the treatment of
inflammation, rheumatoid arthritis, cystitis, post-traumatic
and post ischemic cerebral edema, liver cirrhosis,
Alzheimer's disease, cardiovascular disease, pain, common
cold, allergies, asthma, pancreatitis, burns, virus
infection, head injury, multiple trauma, rhinitis,
hepatorenal failure, diabetes, metastasis, pancreatitis,
30 neovascularization, corneal haze, glaucoma, ocular pain,
ocular hypertension or angio edema, which comprises a
therapeutically effective amount of a compound of Claim 1
and a pharmaceutically acceptable carrier.

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50.

49. A method for the treatment of disease conditions mediated by bradykinin, in a mammalian subject, which comprises administering to said subject a therapeutically effective amount of a compound according to Claim 1.

51.

50. A method for the treatment of inflammation, rheumatoid arthritis, cystitis, post-traumatic and post ischemic cerebral edema, liver cirrhosis, Alzheimer's disease, cardiovascular disease, pain, common cold, allergies, asthma, pancreatitis, burns, virus infection, head injury, multiple trauma, rhinitis, hepatorenal failure, diabetes, metastasis, pancreatitis, neovascularization, corneal haze, glaucoma, ocular pain, ocular hypertension or angio edema, in a mammalian subject, which comprises administering to said subject a therapeutically effective amount of a compound according to Claim 1.

52. 51: A pharmaceutical formulation comprising a compound according to Claim 1, a pharmaceutically acceptable carrier and, optionally, one or more other pharmacologically active ingredients.

20

53.

52. A method of treating, preventing, or ameliorating a disease or condition associated with B1 activity comprising administering to a human or animal subject a therapeutically effective amount of a compound according to Claim 1.

54.

53. The method according to claim 52 wherein the disease or condition is selected from the group consisting of inflammation, inflammatory pain, acute pain, dental pain, back pain, lower back pain, pain from trauma, surgical pain, inflammatory bowel disorders, asthma, and allergic rhinitis.

55. ~~54.~~ The use of a compound according to Claim 1 in the manufacture of a medicament for the treatment of a disease or condition selected from the group consisting of group consisting of inflammation, inflammatory pain, acute pain, 5 dental pain, back pain, lower back pain, pain from trauma, surgical pain, inflammatory bowel disorders, asthma, and allergic rhinitis.

56. ~~55.~~ A compound as in Claim 1 for use in a method of 10 therapeutic treatment for the human or animal body.